

IN THE CLAIMS:

Please cancel without prejudice claims 41, 43, 44, 47-51, 53-55, 57 and 58, such that the claims read as follows:

1. (Previously Presented) A medication delivery apparatus comprising:
 - an antistatic holding chamber comprising a plastic material having a surface resistivity of between about $10E10$ and about $10E12$ ohm/sq, wherein said holding chamber has an input end and an output end spaced apart along a longitudinal axis;
 - a patient interface component connected to said output end and comprising an interior surface defining a flow passage; and
 - a one-way valve disposed adjacent said output end, said one-way valve moveable between an open position and a closed position, wherein said one-way valve has a central opening when in said open position, said central opening defining a flow path along said longitudinal axis, wherein no portion of said interior surface of said patient interface component downstream of said one-way valve intersects said flow path in an orthogonal relationship.
2. (Original) The apparatus of claim 1 wherein said plastic material comprises a polypropylene material.
3. (Previously Presented) The apparatus of claim 1 further comprising a backpiece separate from said holding chamber and comprising an elastomeric material having a surface resistivity of between about $10E10$ and about $10E12$ ohm/sq, wherein said backpiece is connected to said input end of said holding chamber.

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4. (Original) The apparatus of claim 3 wherein said backpiece comprises an opening formed therethrough, said opening shaped and adapted to receive a portion of a pressurized metered dose inhaler.

Claim 5 (Cancelled).

6. (Original) The apparatus of claim 1 wherein said material is selected from the group consisting of polypropylene, polycarbonate, polystyrene, nylon, acrylonitrile butadiene styrene, high density polyethylene, acetal, polybutylene terephthalate, and polyethylene terephthalate glycol.

7. (Original) The apparatus of claim 1 wherein at least a portion of said holding chamber is see-through.

8. (Original) The apparatus of claim 1 wherein said surface resistivity of said plastic material is between about $10E10$ and about $10E11$ ohm/sq.

9. (Previously Presented) The apparatus of claim 1 further comprising a second antistatic component separate from said holding chamber and comprising a material having a surface resistivity of between about $10E10$ and about $10E12$ ohm/sq, and wherein said second antistatic component is connected to said holding chamber.

10. (Previously Presented) The apparatus of claim 9 wherein said second antistatic component comprises said patient interface component connected to said output end of said holding chamber.

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11. (Previously Presented) The apparatus of claim 9 wherein said second antistatic component comprises a backpiece connected to said input end of said holding chamber.

12. (Original) The apparatus of claim 11 wherein said backpiece comprises an elastomeric material.

Claims 13-16 (Cancelled).

17. (Original) The apparatus of claim 11 wherein said backpiece comprises an opening formed therethrough, said opening shaped and adapted to receive a portion of a pressurized metered dose inhaler.

Claim 18 (cancelled).

19. (Original) The apparatus of claim 11 wherein said material comprises a thermoplastic elastomer material.

20. (Previously Presented) The apparatus of claim 9 wherein said material of said second antistatic component is selected from the group consisting of a polyurethane elastomer, polyester elastomer, styrenic elastomer and olefinic elastomer.

21. (Previously Presented) The apparatus of claim 9 wherein at least a portion of said holding chamber and said second antistatic component is see-through.

Claims 22-58 (Cancelled).

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59. (Previously Presented) The apparatus of claim 1 wherein said patient interface component comprises a mouthpiece.

60. (Previously Presented) The apparatus of claim 59 wherein said interior surface of said mouthpiece is not antistatic.